

In the Claims

1. (amended) A drug delivery composition for nasal administration comprising ICAM-1 and a bioadhesive material, wherein the bioadhesive material is in a liquid formulation comprising a polymeric material, wherein the ICAM-1 is present in the liquid formulation in a concentration between about 0.01 and 20% by weight per volume, and wherein the composition delivers to the nasal cavity an antivirally effective amount of ICAM-1.

2. (Amended) The drug delivery composition according to claim 1 wherein the bioadhesive material is a chitosan solution.

3. (Amended) The drug delivery composition according to claim 2 wherein the chitosan is in the solution in a concentration in the range of 0.2 - 2.0% w/v.

4. (Amended) The [A] drug delivery composition according to claim 2 [or 3] wherein the ICAM-1 is present in the chitosan solution in a concentration in the range of 0.2 to 5% w/v.

5. (Amended) A drug delivery composition for nasal administration comprising ICAM-1 and a [according to claim 1 wherein the] bioadhesive material in a dry powder formulation, wherein the bioadhesive material is a plurality of microspheres made from a material selected from the group consisting of starch, chitosan, gelatin, hyaluronic acid, alginate, and gellan, wherein the ICAM-1 content of the formulation is between about 0.1 and 50% by weight, and wherein the composition delivers to the nasal cavity an antivirally effective amount of ICAM-1.

Please cancel claim 6.

7. (Amended) The [A] drug delivery composition according to claim 5 [or 6]
wherein the ICAM-1 is present in an amount of 1% to 20% w/w of the microspheres.

Please cancel claim 8.

9. (Amended) The [A] drug delivery composition according to claim 1
wherein the polymeric material is selected from the group consisting of gellan gum,
alginate, welan, xanthan, [or] and rhamsan.

10. (Amended) The [A] drug delivery composition according to claim [8 or 9]
1 wherein the polymeric material is provided in a concentration of 0.1% to 5% w/v.

11. (Amended) [A] The drug delivery composition according to [any one of
claims 9-10] claim 8 wherein the ICAM-1 is present in the formulation in an amount of
0.2% to 5% w/v.

12. (Amended) A method of delivering ICAM-1 to the nasal cavity to
increase its effectiveness therein comprising
administering the ICAM-1 in a drug delivery composition additionally comprising
a bioadhesive material, wherein the bioadhesive material is in a liquid formulation
comprising a polymeric material or is in a dry powder formulation comprising a plurality
of microspheres made from a material selected from the group consisting of starch,
chitosan, gelatin, hyaluronic acid, alginate, and gellan, and wherein the composition
delivers to the nasal cavity an antivirally effective amount of ICAM-1.

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Favorable consideration is earnestly solicited.

Respectfully submitted,

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